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**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**

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of

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**Complete if Known**

Application Number	10/661,780
Filing Date	09/15/2003
First Named Inventor	PHILIPPE BOUCHARD
Art Unit	1614
Examiner Name	Brian Yong S. Kwon
Attorney Docket Number	098501-0305998

**U. S. PATENT DOCUMENTS**

Examiner Initials*	Cite No. <sup>1</sup>	Document Number Number-Kind Code <sup>2</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		US- 6,077,523	06/20/2000	Romano Deghenghi	
		US- 6,022,860	02/08/2000	Engel et al.	
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**FOREIGN PATENT DOCUMENTS**

Examiner Initials*	Cite No. <sup>1</sup>	Foreign Patent Document Country Code <sup>3</sup> Number <sup>4</sup> Kind Code <sup>5</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>

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# INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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**Complete if Known**

Application Number

10/661,780

Filing Date

09/15/2003

First Named Inventor

PHILIPPE BOUCHARD

Art Unit

1614

Examiner Name

Brian Yong S. Kwon

Attorney Docket Number

098501-0305998

**NON PATENT LITERATURE DOCUMENTS**

Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
		NESTOR, JR. ET AL., "Potent gonadotropin releasing hormone antagonists with low histamine-releasing activity," J. Med. Chem., Institute of Bio-Organic Chemistry, Vol. 35 ( No. 21), p. 3942-3948, (October 16, 1992).	
		DEGHENGHI ET AL., "Antarelix (EP 24332) a novel water soluble LHRH antagonist," Biomed. Pharmacother., Europeptides (Argenteuil, France), Vol. 47 ( No. 2-3), p. 107-110, (1993).	
		RIVIER ET AL., "Gonadotropin-releasing hormone antagonists with N omega-triazolylornithine, -lysine, or -p-aminophenylalanine residues at positions 5 and 6," J. Med. Chem., Salk Institute for Biological Studies (La Jolla, California ), Vol. 35 ( No. 23), p. 4270-4278, (November 13, 1992).	
	See Attachment 1		
		HAVIV ET AL., "In vitro and in vivo activities of reduced-size antagonists of luteinizing hormone-releasing hormone," J. Med. Chem., TAP Pharmaceuticals, Inc. (Abbott Park, IL), Vol. 37 ( No. 5), p. 701-705, (March 4, 1994).	
		WEINBAUER ET AL., "Comparison of the antgonadotropic activity of three GnRH antagonists (Nal-Glu, Antide and Cetrorelix) in a non-human primate model (Macaca fascicularis)," Andrologia, Institute of Reproductive Medicine of the University (Munster, Germany), Vol. 25 ( No. 3), p. 141-147, (May-June 1993).	

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<sup>1</sup> Applicant's unique citation designation number (optional). <sup>2</sup> Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. **SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

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## Addendum

Page 3 of 3

Attachment 1

- a.
- b. STOECKEMANN ET AL., "Effects of the luteinizing-hormone-releasing hormone (LHRH) antagonist ramorelix (hoe013) and the LHRH agonist buserelin or dimethylbenz[*a*]anthracene-induced mammary carcinoma: studies with slow-release formulations," J. Cancer Res. Clin. Oncol., Hoechst AG, Pharma-Research (Frankfurt/Main, Germany), Vol. 119 ( No. 8), p. 457-462, (1993).
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